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- (iv) Written approval from the Administrator of the fuel specifications was provided prior to the start of testing.
- (5) The specification range of the fuels to be used under paragraphs (b) (2), (3), and (4) of this section shall be reported in accordance with §86.082–21(b)(3).
- (c) Fuels not meeting the specifications set forth in this section may be used only with the advance approval of the Administrator.

[46 FR 50494, Oct. 13, 1981, and 47 FR 49807, Nov. 2, 1982]

§86.308-79 Gas specifications.

- (a) Analyzer gases. (1) Calibration gases for the CO and CO_2 analyzers shall have zero grade nitrogen as a diluent. Combined CO and CO_2 span gases are permitted. Zero grade nitrogen shall be the diluent for CO and CO_2 span gases.
- (2) Calibration or span gases for the hydrocarbon analyzer shall be propane with zero-grade nitrogen as a diluent when testing gasoline-fueled engines. For Diesel engine tests the diluent shall be zero-grade air.
- (3) Calibration or span gases for the $NO_{\rm X}$ analyzer shall be NO named as $NO_{\rm X}$ with a maximum NO_2 concentration of 5 percent of the nominal value. Zero-grade nitrogen shall be the diluent.
- (4) Zero-grade gases for hydrocarbon analyzers shall be nitrogen when testing gasoline-fueled engines and air when testing Diesel engines.
- (5) Zero-grade gases for the carbon monoxide, carbon dioxide and oxides of nitrogen analyzers shall be either zero-grade air or zero-grade nitrogen.
- (6) The allowable zero grade gas (air or nitrogen) impurity concentrations shall not exceed 2 ppmC hydrocarbon, 10 ppm carbon monoxide, 400 ppm carbon dioxide and 1 ppm nitric oxide.
- (7) "Zero-grade air" includes artificial "air" consisting of a blend of nitrogen and oxygen with oxygen concentrations between 18 and 21 mole percent.
- (b) Calibration gas. Calibration gas values are to be derived from NBS "Standard Reference Materials" (SRM's) or other gas standards approved by the Administrator. The un-

- certainty of the assigned calibration gas values shall not exceed 2.0 percent of the assigned value. The uncertainty is defined as the sum of the precision errors (at the 90 percent confidence level) and the bias errors. Precision and bias errors apply to both the equipment and the derivation procedures.
- (c) Span gas. Span gas values are to be derived from NBS "Standard Reference Materials" (SRM's) or other gas standards approved by the Administrator. The uncertainty of the assigned calibration gas values shall not exceed 3.0 percent of the assigned value. The uncertainty is defined as the sum of the precision errors (at the 90 percent confidence level) and the bias errors. Precision and bias errors apply to both the equipment and the derivation procedures.
- (d) Hydrocarbon analyzer fuel—(1) The fuel shall contain 40 ±2 percent hydrogen. The balance shall be helium. The mixture shall contain less than 2 ppmC hydrocarbon.
- (2) Alternate pure hydrogen fuel. Some HFID's are designed to operate on pure hydrogen. Generally the HFID fuel is incompatible with good relative hydrocarbon response.
- (i) For Diesel engines this fuel is not recommended. However, this fuel may be used if the engine manufacturer demonstrates, on each basic combustion system (i.e., 4 cycle DI, 2 cycle DI, 4 cycle pre-cup, etc.) that an HFID using this fuel produces comparable results to an HFID using 40% H₂/60% He fuel. These data must be submitted to and approved by the Administrator prior to testing. Pure H₂ fuel, that may be allowed for testing, must contain at least 99.0 percent hydrogen and contain less than 2 ppmC hydrocarbon.
- (ii) For gasoline-fueled engines, pure hydrogen fuel for the HFID is not allowed.
- (e) Hydrocarbon analyzer burner air. The concentration of oxygen must be within 1 mole percent of the oxygen concentration of the burner air used in the latest oxygen interference check (%O₂ I). If the difference in oxygen concentration is greater than 1 mole percent, then the oxygen interference must be checked and the analyzer adjusted if necessary, to meet the %O₂ I

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requirements. The burner air must contain less than 2 ppmC hydrocarbon.

(f) Oxygen interference check gases shall contain propane with 350 ppmC ±75 ppmC hydrocarbon. The concentration value shall be determined to calibration gas tolerances by chromatographic analysis of total hydrocarbons plus impurities or by dynamic blending. Nitrogen shall be the predominant diluent with the balance oxygen. Blends required for gasolinefueled and Diesel engine testing are as follows:

Applicability	O ₂ concentration (percent)	Balance
Diesel Diesel and gasoline Do Gasoline	21 (20 to 22)	$\begin{array}{c} N_2 \\ N_2 \\ N_2 \\ N_2 \end{array}$

(g) Proportioning and blending devices may be used to obtain required gas concentration.

[42 FR 45154, Sept. 8, 1977, as amended at 46 FR 50495, Oct. 13, 1981; 47 FR 49807, Nov. 2, 1982]

§86.309-79 Sampling and analytical system; schematic drawing.

(a) Any variation from the specifications in this subpart including performance specifications and emission detection methods may be used only with prior approval by the Administrator.

(b) Schematic drawing. (1) An example of a sampling and analytical system which may be used for testing under this subpart is shown in Figure D79–1. All components or parts of components that are wetted by the sample or corrosive calibration gases shall be either chemically cleaned stainless steel or inert material, e.g. polytetrafluoroethylene resin. The use of "gauge savers" or "protectors" with nonreactive diaphragms to reduce dead volumes is

permitted. The specific detection methods to be used for each exhaust component can be found in §86.316 for CO. §86.317 for HC, and §86.318 for NO_X. Additional components such as instruments, valves, solenoids, pumps, switches, etc. may be employed to provide additional information and coordinate the functions of the component systems.

- (2) The following requirements must be incorporated in each system used for testing under this subpart.
- (i) All analyzers must obtain the sample to be analyzed from the same sample line.
- (ii) The sample transport system from the engine exhaust pipe to the HC analyzer and the NO_X analyzer must be heated as is indicated in Figure D79-1.
- (iii) Carbon monoxide and carbon dioxide measurements must be made on a dry basis. Specific requirements for the means of drying the sample can be found in §86.309 (b)(5) and §86.311(e).
- (iv) All NDIR analyzers must have a pressure gauge immediately downstream of the analyzer. The gauge tap must be within 2 inches of the analyzer exit port. Gauge specifications can be found in §86.309(b)(3).
- (v) All bypass and analyzer flows exiting the analysis system must be measured. Capillary flows such as in HFID and CL analyzers are excluded. For each ND IR analyzer with a flow meter located upstream of the analyzer, an upstream pressure gauge must be used. The gauge tap must be within 2 inches of the analyzer entrance port.
- (vi) Calibration or span gases for the NO_X measurement system must pass through the NO_2 to NO converter.
- (vii) The temperature of the NO₂ to NO converter must be displayed continuously.